Cardiovascular Ultrasound



Open Access Research

A comparison of Power Doppler with conventional sonographic imaging for the evaluation of renal artery stenosis

Agatino Manganaro, Giuseppe Ando'*, Antonino Salvo, Andrea Consolo, Frank Coppolino and Domenico Giannino

Address: Dipartimento Clinico - Sperimentale di Medicina e Farmacologia, Università degli Studi di Messina, Policlinico "G. Martino", Messina,

Email: Agatino Manganaro - manganaro51@unime.it; Giuseppe Ando'* - giuseppe.ando@tiscali.it; Antonino Salvo - giuseppe.ando@tiscali.it; Andrea Consolo - giuseppe.ando@tiscali.it; Frank Coppolino - giuseppe.ando@tiscali.it; Domenico Giannino - giuseppe.ando@tiscali.it

* Corresponding author

Published: 14 January 2004 Cardiovascular Ultrasound 2004, 2:1

Accepted: 14 January 2004 This article is available from: http://www.cardiovascularultrasound.com/content/2/1/1

© 2004 Manganaro et al; licensee BioMed Central Ltd. This is an Open Access article: verbatim copying and redistribution of this article are permitted in all media for any purpose, provided this notice is preserved along with the article's original URL.

Received: 18 December 2003

Abstract

Background: Power Doppler (PD) has improved diagnostic capabilities of vascular sonography, mainly because it is independent from the angle of insonation. We evaluated this technique in a prospective comparison with conventional imaging, consisting in Duplex and Color Doppler, for the evaluation of Renal Artery (RA) stenosis.

Methods: Sensitivity, specificity and predictive values of PD and conventional imaging were assessed in a blinded fashion on eighteen patients, 9 with angiographic evidence of unilateral RA stenosis (hypertensive patients) and 9 with angiographically normal arteries (control group). PD images were interpreted with an angiography-like criteria.

Results: In the control group both techniques allowed correct visualization of 16 out of the 18 normal arteries (93% specificity). Only in five hypertensive patients RA stenosis was correctly identified with conventional technique (56% sensitivity and 86% negative predictive value); PD was successful in all hypertensive patients (100% sensitivity and negative predictive value), since the operators could obtain in each case of RA stenosis a sharp color signal of the whole vessel with a clear "minus" at the point of narrowing of the lumen. All results were statistically significant (p < 0.01).

Conclusions: This study demonstrates that PD is superior to conventional imaging, in terms of sensitivity and specificity, for the diagnosis of RA stenosis, because it allows a clear visualization of the whole stenotic vascular lumen. Especially if it is used in concert with the other sonographic techniques, PD can enable a more accurate imaging of renovascular disease with results that seem comparable to selective angiography.

Background

Duplex/CD sonography is used as a routine method for the assessment of RAs [1,2]. This technique is a reliable diagnostic tool in order to display vascular structures as well as velocity and flow direction of red blood cells by

mapping the Doppler frequency shift information. However, the detection of RAs through the anterior abdominal wall with this conventional technique is hampered in a substantial group of patients either by anatomical origin of RAs from a plane that may be almost perpendicular to the ultrasonic beam [3-5], or by several technical issues like bowel gas, breathing movements and obesity. In such cases RAs exploration through modified windows, like the margino-renal, lateral window, is crucial [2]. Furthermore, it is intriguing to complete the sonographic evaluation with PD. PD generates an intravascular color map reflecting the integrated power in the Doppler signal, which essentially depends on the density of red blood cells within the sample volume. Since increased sensitivity to visualize the continuity of blood flow in arterial stenosis is claimed for PD, we evaluated this new technique in a prospective comparison with conventional imaging for the diagnosis of RA stenosis.

Methods Population

For the purpose of this study, we considered nine patients (6 men; mean age 62, range 48-76 years) referred from January 2002 to October 2003 for evaluation of hypertension. In particular, these patients had a clinical suspicion of RA stenosis, because of hypertension resistant to the combination of different antihypertensive drugs (betablockers, vasodilators and diuretics) associated either with moderate renal failure or with carotid and peripheral atherosclerotic disease. With selective renal angiography, we demonstrated in these patients significant unilateral (5 cases of right, 4 cases of left) RA stenosis; they were considered as the group of hypertensive patients. As a control group, we enrolled nine patients out of the population undergoing left heart catheterization for coronary angiography (5 men; mean age 66; range 55-74 years), matched for age and other risk factors, in whom we demonstrated angiographically normal RAs. All patients gave their written informed consent to participate in the study, that was aimed at comparing the results of conventional technique to those obtained with PD in the identification and characterization of RA stenosis.

The sonographic evaluation of RAs of the hypertensive patients had been performed the day before angiography for obvious clinical reasons, whereas controls underwent a blinded evaluation two days after coronary angiography. Such examinations have been carried out by three independent physicians, all having at least a five-year-experience in vascular sonography.

Imaging technique

Renal sonography is performed in our Department with ATL machines (Advanced Technology Laboratories, Bothell, Washington, USA): "Ultramark 9", "Apogee 800" and "HDI 5000". The last two are equipped with PD facilities and have been used for the present study.

For the diagnosis of RA stenosis with the conventional technique, we perform a direct imaging of the main RA,

with the color window used to look for the aliasing phenomenon in order to sample the area of the stenosis with PW Doppler interrogation, as widely validated in several previous studies [2,6]. We consider 200 cm/sec as a diagnostic cutoff for significant RA stenosis. When performing PD examination, we give an estimation of the degree of RA stenosis from the color window itself, considering color as a contrast media and giving an angiography-like interpretation of the images. Furthermore, we use the color window to place the sample volume at the point of the assumed stenosis for a supplemental PW interrogation.

The patient undergoing renal sonography is kept recumbent in a lateral position: a left decubitus to explore the right RA and a right decubitus to explore the left RA. A short-axis scanning of the aorta has to be obtained, in order to align as much as possible the ultrasonic beam to the axis of RA, resulting in a low angle of attack. Thus, our purpose is to obtain the highest signal intensity of the RA (Figure 1) using an orthogonal variant of the "banana peel" view [2].

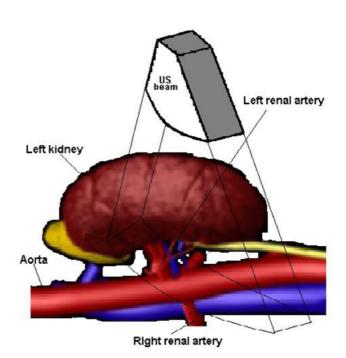


Figure I
Short axis approach from the lateral window for left
RA imaging. The patient is kept in a right decubitus to
explore the left RA. A short-axis scanning of the aorta is
obtained, in order to align as much as possible the ultrasonic
beam to the axis of the left RA.

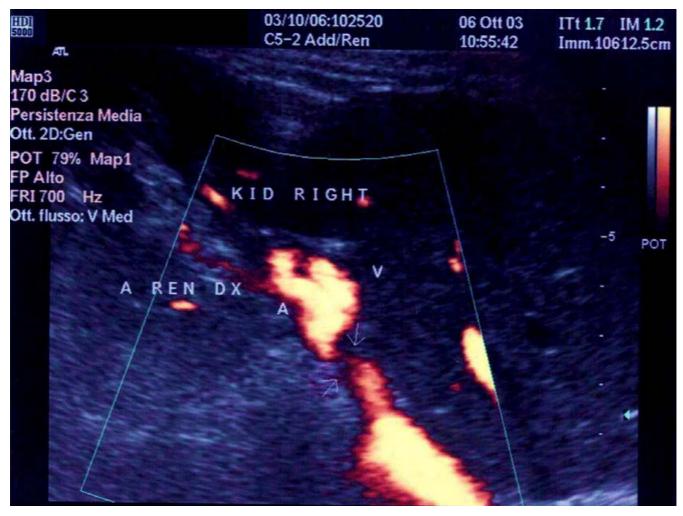


Figure 2

Power Doppler image of a stenosis of right RA. This image has been obtained from one of the patients described in the text with the ATL HDI 5000 machine. The arrows indicate the stenosis. The distal lumen of the RA is normal. KID, kidney. A REN DX, right renal artery. V, vein. A, artery.

Statistical analysis

Sensitivity, specificity and predictive values, with 95% CI, of both PD and conventional imaging have been determined on the thirty-six RAs that were examined by using a two-sided Fisher's exact test, with an alpha level of 0,05.

Results

Control group

both conventional technique and PD allowed a clear visualization and a sharp spectral analysis of 16 out of the 18 angiographically normal RAs that were studied (93% specificity of both techniques). In two very obese women, the visualization of the left RA was inadequate both with CD and PD: consequently, placement of the PW sample volume and velocity measurement were impossible. For the

purpose of the statistical analysis, these two cases were considered as false positives.

Hypertensive patients

five cases (4 right, 1 left) of RA stenosis were correctly diagnosed with the conventional technique (56% sensitivity and 86% negative predictive value); in all these cases a PSV higher than 200 cm/s was recorded when the sample volume was placed at the aliasing point of the CD images. In the PD images of these narrowed RAs, color signal clearly contrasted a substantially narrowed vascular lumen in the areas of the ostia; the middle and the distal lumen were normal in all patients, both with CD and PD. Both operators were in agreement that, according to such PD images, the degree of stenosis had to be more than

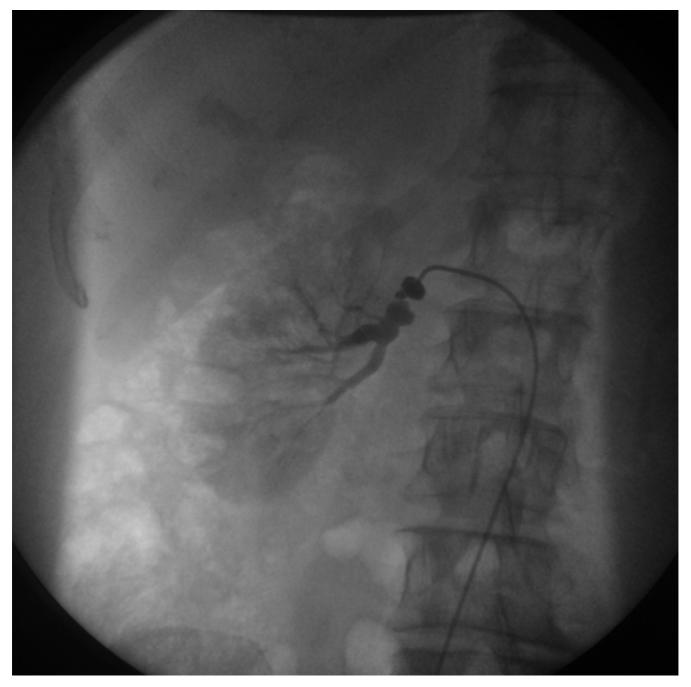


Figure 3
Selective angiography of a stenosis of right RA. This image has been obtained from the same patient of figure 2 at the time of percutaneous angioplasty. The stenosis of the right RA is evident.

70%. In the other four cases (1 right, 3 left) of RA stenosis, the conventional sonographic evaluation had failed to clearly demonstrate an obstructive pathology of RAs, both through the anterior abdominal wall and the flank. RAs scanning had been extremely demanding because of sub-

optimal technical conditions and the diagnosis of RA stenosis had not been possible because direct criteria had not been met; in all patients in whom an almost stable PW interrogation was possible, the PSV was lower than 180 cm/s. The images obtained with PD were similar to those

	Power Doppler	95% CI	Conventional imaging	95% CI
Sensitivity	100%	66%–100%	56%	21%–86%
Specificity	93%	76%–99%	93%	76%–99%
Positive Predictive Value	82%	48%–98%	71%	29%–96%
Negative Predictive Value	100%	86%-100%	86%	68%–96%
p (Fisher's exact test)	< 0.0001		0,0056	

Table: Overall sensitivity, specificity and predictive values of Power Doppler and conventional imaging.

obtained in the former patients, with the color signal clearly in contrast (not only upstream but even downstream) with a significant narrowing of the vascular lumen, thus permitting the operators to advance the diagnosis of RA stenosis. Neither the use of PD color map to place the sample volume for PW interrogation at the site of the stenosis permitted to demonstrate a cutoff PSV, because of the same lack of stability of the PW signal. The degree of stenosis that had been demonstrated with angiography seemed in all cases comparable to that estimated with PD imaging (Figure 2 and 3).

Overall data are summarized in the table 1. The four patients that underwent angioplasty with stent implantation had a successful outcome, with almost complete resolution of hypertension and no clinical need for further sonographic evaluations.

Discussion

RA stenosis is a common cause of secondary hypertension and angiography is considered to be the gold standard for the morphologic visualization of RA stenosis [7,8]. Yet, angiography is costly, invasive, and associated with inherent morbidity. Therefore, it cannot be used as a screening method and it must be reserved for the confirmation of a diagnosis suggested by other techniques and for interventional procedures. Either computed tomographic or magnetic resonance angiography and captopril renography can evaluate the renal perfusion and the haemodynamic significance of a stenosis [1,9,10]. All these techniques are safer alternatives to angiography; nevertheless, scintigraphy does not provide any anatomical information and, moreover, high cost and not widespread availability of all of them may somehow preclude a routine use in clinical practice [11].

Sonography, as a combination of grey-scale B-mode, CD and PW spectral Doppler, is a totally non-invasive technique that is widely used as a first-line examination to advance the diagnosis of RA stenosis [1,2,6]. Yet, conventional sonography often requires a high degree of technical expertise in order to provide accurate results. In other words, it may be technically very demanding and time

consuming to clearly identify and interrogate RAs because of their angle of origin from the aorta and their sinuous course; further issues arise from anatomical variants, that can result in false negatives, and from technically inadequate test conditions such as body habitus, inability to hold the breath, bowel gas.

As an alternative and indirect method of identifying RA stenosis, many groups use to analyze intrarenal waveforms, as pointed out by the original observations by Handa [12] and Stavros [13], looking for the tardus-parvus effect. Good results can be obtained only with some quantitative parameters [12-14], whose value in clinical practice has been reviewed elsewhere [2]. Although such "indirect" parameters can overcome technical demands inherent to main RAs interrogation, many groups -as we do- consider them less reliable than "direct" ones [15-18]. Indeed, intrarenal PW examination may be relatively easy in patients without abnormalities, as our control group. To obtain diagnostic-quality spectral signals when the clinical suspicion is high, however, is almost as challenging and time consuming as to obtain satisfactory images of the main RAs with CD and PW interrogation. Moreover, the tardus-parvus technique is limited by a low sensitivity due to the intrinsic variability of Doppler waveforms in patients without abnormalities [16-18] and by the fact that the degree of stenosis necessary to consistently cause downstream flow changes is still unknown: the simple look at the waveform shape and the observation of the presence or absence of an end-systolic peak may only allow to differentiate normal from generically abnormal flow [1,2].

The development of PD has been permitting further diagnostic possibilities for renal sonography [19,20]. CD and PD have distinct applications: the knowledge of advantages and limitations of each is essential for their proper application. Both CD and PD elaborate the Doppler frequency shift. The main difference consists in the parameters of the signal that are processed: CD mode processes the mean frequency shift to provide color intensity representing velocity, and the phase shift to indicate flow direction. PD mode generates an intravascular color

map reflecting the integrated power in the Doppler signal [21]. Such a parameter depends on the amount of red blood cells producing the Doppler frequency shift, regardless of their flow velocity and direction.

Providing a representation of a different property of blood flow, PD has several advantages over CD; it has greater sensitivity to detect the blood flow itself and, with the same angle of insonation, PD images permit better definition of the intravascular surfaces and visualization of continuity of flow than CD ones. In other words, PD images are independent from the angle of insonation [21]. This is particularly useful in arterial stenosis, since PD imaging results in an angiography-like visualization of the whole vascular lumen [22,23]. Indeed, PD permits to clearly represent in color the low-velocity post-stenotic flow. This could be even possible with broad band CD flow, keeping PRF as low as possible; with such a modification of PRF, yet, the aliasing phenomenon would lose its significance and the frame rate would be extremely low. Moreover, PD improves the visibility of vessels that are sinuous or kinked or lying on a plane perpendicular to the ultrasonic beam, due to its independence from the angle of insonation. Yet, PD disadvantages mainly consist in long scanning time that makes PD images susceptible to soft-tissue flash artifacts and movement artifacts; moreover, PD does not yield directional or velocity information and, consequently, does not permit to detect aliasing, although it gives the operator the possibility to place the PW sample volume at the most stenotic tract of the vessel. Finally, the frame rate is much lower with PD than with CD, so that arteries are not distinguished from veins on the basis of their color pulsatility. Accordingly, PD is a useful tool, but it is complementary to Duplex/CD in vascular sonography: they should be used together for optimal diagnostic results, CD and aliasing being the guides to PD activation.

Although there are no universally accepted criteria to make a diagnosis of RA stenosis with PD, displayed perfusion characteristics are similar to those obtained from angiography (Figure 2 and 3). According to our observations, therefore, if an intriguing, angiography-like, interpretation of PD images is given, even of a single frame with a color "minus", this technique is likely to permit an estimation, though not as exact as with angiography [20,24], of the degree of stenosis, thus of the haemodynamic significance of a plaque. Furthermore, with the appropriate learning curve, PD could shorten the overall imaging time when the angiography-like diagnosis of stenosis is obtained, mainly because it would be not necessary to perform long scans in deep inspiration in order to achieve a stable PW flow trace.

In our experience, the evaluation through the lateral window seems particularly useful in the case of left RA steno-

sis: this is due, in part, to the absence of the good sonographic window created by the liver for right RA. Radiological studies have showed that the left RA has in most cases a lateral origin from the aorta [3-5]: thus the lateral approach, with the patient recumbent on the right side, allows the operator to avoid usual abdominal obstacles and makes the ultrasonic beam aligned as much as possible to the axis of the vessel: the result is the highest intensity of the left RA Doppler signal.

The main limitation of this study is the small number of patients the we could enroll, due to the low prevalence of RA stenosis in the whole population of patients with hypertension. Moreover, the overall high specificity (93%) of sonography is not surprising and is consistent with the literature [1]. In our hypertensive patients there was a relevant clinical suspicion of RA stenosis, quite apart from the subsequent angiographic demonstration: this circumstance represents a relevant bias in the selection of patients to be investigated, as to let us obtain high sensitivity and negative predictive values, especially with PD (table 1). Therefore, a larger prospective study is needed, in order to compare PD-guided sonography with conventional sonography for the diagnosis of RA stenosis in an unselected cohort of patients with hypertension.

Conclusions

This study demonstrates that PD is superior to conventional imaging, in terms of sensitivity and specificity, for the diagnosis of RA stenosis, because it allows a clear visualization of the whole stenotic vascular lumen. Due to its intrinsic limitations, PD cannot replace conventional sonographic techniques and especially CD, whose aliasing phenomenon is the most useful guide to activate PD during the examination. When PD is used as an adjunctive tool in vascular sonography, it enables a more accurate imaging of renovascular disease with results that seem comparable to selective angiography.

List of abbreviations

CD Color Doppler

CI Confidence Interval

PD Power Doppler

PRF Pulse Repetition Frequency

PSV Peak Systolic Velocity

PW Pulsed Wave

RA Renal Artery

RAs Renal Arteries

Competing interests

None declared.

Authors' contributions

AM and GA designed the study, AM, DG and AS carried out the sonographic evaluations, AC and GA performed the angiographic examinations.

GA performed literature research and prepared, edited and reviewed the manuscript.

FC and DG participated in the design and coordination of the study, and AM, GA, AC and AS are guarantors of its integrity.

All authors read and approved the final manuscript.

References

- Rankin SC, Saunders AJ, Cook GJ, Scoble JE: Renovascular disease. Clin Radiol 2000, 55:1-12.
- Lee H, Grant EG: Sonography in renovascular hypertension. J Ultrasound Med 2002, 21:431-441.
- Verschuyl EJ, Kaatee R, Beek FJ, Pasterkamp G, Bush WH, Beutler JJ, van der Ven PJ, Mali WP: Renal artery origins: location and distribution in the transverse plane at CT. Radiology 1997, 203:71-75.
- Beregi JP, Mauroy B, Willoteaux S, Mounier-Vehier C, Remy-Jardin M, Francke J: Anatomic variation in the origin of the main renal arteries: spiral CTA evaluation. Eur Radiol 1999, 9:1330-1334.
- Garcier JM, De Fraissinette B, Filaire M, Gayard P, Therre T, Ravel A, Boyer L: Origin and initial course of the renal arteries: a radiological study. Surg Radiol Anat 2001. 23:51-55.
- ological study. Surg Radiol Anat 2001, 23:51-55.

 Hua HT, Hood DB, Jensen CC, Hanks SE, Weaver FA: The use of colorflow duplex scanning to detect significant renal artery stenosis. Ann Vasc Surg 2000, 14:118-124.
- Safian RD, Textor SC: Renal-artery stenosis. N Engl J Med 2001, 344:431-442.
- Haller H: Arteriosclerotic renal artery stenosis: conservative versus interventional management. Heart 2002, 88:193-197.
- Bongers V, Bakker J, Beutler JJ, Beek FJ, De Klerk JM: Assessment of renal artery stenosis: comparison of captopril renography and gadolinium-enhanced breath-hold MR angiography. Clin Radiol 2000, 55:346-353.
- De Cobelli F, Venturini M, Vanzulli A, Sironi S, Salvioni M, Angeli E, Scifo P, Garancini MP, Quartagno R, Bianchi G, Del Maschio A: Renal arterial stenosis: prospective comparison of color Doppler US and breath-hold, three-dimensional, dynamic, gadolinium-enhanced MR angiography. Radiology 2000, 214:373-80.
- Nelemans PJ, Kessels AG, De Leeuw P, De Haan M, van Engelshoven J: The cost-effectiveness of the diagnosis of renal artery stenosis. Eur J Radiol 1998, 27:95-107.
- Handa N, Fukunaga R, Etani H, Yoneda S, Kimura K, Kamada T: Efficacy of echo-Doppler examination for the evaluation of renovascular disease. Ultrasound Med Biol 1988, 14:1-5.
- 13. Stavros AT, Parker SH, Yakes WF, Chantelois AE, Burke BJ, Meyers PR, Schenck JJ: Segmental stenosis of the renal artery: pattern recognition of tardus and parvus abnormalities with duplex sonography. Radiology 1992, 184:487-492.
- Riehl J, Schmitt H, Bongartz D, Bergmann D, Sieberth HG: Renal artery stenosis: evaluation with colour duplex ultrasonography. Nephrol Dial Transplant 1997, 12:1608-1614.
- Richardson D, Foster J, Davison AM, Irrving HC: Parvus tardus waveform suggesting renal artery stenosis-remember the more proximal stenosis. Nephrol Dial Transplant 2000, 15:539-543.
- Kliewer MA, Tupler RH, Carroll BA, Paine SS, Kriegshauser JS, Hertzberg BS, Svetkey LP: Renal artery stenosis: analysis of Doppler waveform parameters and tardus-parvus pattern. Radiology 1993, 189:779-787.
- Baxter GM, Aitchison F, Sheppard D, Moss JG, McLeod MJ, Harden PN, Love JG, Robertson M, Taylor G: Colour Doppler ultrasound

- in renal artery stenosis: intrarenal waveform analysis. Br J Radiol 1996, 69:810-815.
- Motew SJ, Cherr GS, Craven TE, Travis JA, Wong JM, Reavis SW, Hansen KJ: Renal duplex sonography: main renal artery versus hilar analysis. J Vasc Surg 2000, 32:462-471.
- Chen P, Maklad N, Redwine M: Color and Power Doppler imaging of the kidneys. World | Urol 1998, 16:41-45.
- Helenon O, Correas JM, Chabriais J, Boyer JC, Melki P, Moreau JF: Renal vascular Doppler imaging: clinical benefits of power mode. Radiographics 1998, 18:1441-1454.
- Rubin JM, Bude RO, Carson PL, Bree RL, Adler RS: Power Doppler US: a potentially useful alternative to mean frequency-based color Doppler US. Radiology 1994, 190:853-856.
- Steinke W, Ries S, Artemis N, Schwartz A, Hennerici M: Power Doppler imaging of carotid artery stenosis. Comparison with color Doppler flow imaging and angiography. Stroke 1997, 28:1981-1987.
- 23. Bluth El, Sunshine JH, Lyons JB, Beam CA, Troxclair LA, Althans-Kopecky L, Crewson PE, Sullivan MA, Smetherman DH, Heidenreich PA, Neiman HL, Burkhardt JH: Power Doppler imaging: initial evaluation as a screening examination for carotid artery stenosis. *Radiology* 2000, 215:791-800.
- Claudon M, Winninger D, Briancon S, Pesque P: Power Doppler US: evaluation of the morphology of stenoses with a flow phantom. Radiology 2001, 218:109-117.

Publish with **Bio Med Central** and every scientist can read your work free of charge

"BioMed Central will be the most significant development for disseminating the results of biomedical research in our lifetime."

Sir Paul Nurse, Cancer Research UK

Your research papers will be:

- available free of charge to the entire biomedical community
- peer reviewed and published immediately upon acceptance
- cited in PubMed and archived on PubMed Central
- yours you keep the copyright

Submit your manuscript here: http://www.biomedcentral.com/info/publishing_adv.asp

